

TRISHEVSKIY, I.S.; GAMERSHTEYN, V.A.; SKOKOV, F.I.; AKIMOV, E.P.

Dependence of metal hardening on the conditions of shaping  
and the width of the initial ingot. Sber. trud. UNIIM  
no. 7.1:208-215 '65. (MIRA 18:11)

TRISHEVSKIY, I.S.; STUKALOV, V.P.; SKOKOV, F.I., DRAPIKO, P.Ye.

Developing and studying the technology of producing rolled shapes with elements bent to 180°. Sbor.trud. UNIM no.11:216-231 '65. (MIRA 18:11)

PN25

S/183/62/000/002/012/013  
B102/B108

49,3300

AUTHOR: Skokov, I. V.

TITLE: Comparison of the sensitivities of a double-ray and a multi-ray interferometer for measuring small variations in refractive index

PERIODICAL: Moscow. Universitet. Vestnik. Seriya III. Fizika, astronomiya, no. 2, 1962, 82-87

TEXT: The author calculated the sensitivities of a double-ray and a multi-ray interferometer used for measuring small variations in refractive index  $n$  of gases. The sensitivities depend on the length of the rays and on the accuracy of recording the changes in the interference pattern. The latter are assumed to be recorded by a photometric method, the dimensions of the test object being equal in both cases; the photometric errors are assumed to be equal and the interferometers to be hit by parallel rays of monochromatic light. External effects are ignored. The sensitivities are calculated from the intensity distributions

Card 1/2

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Comparison of the sensitivities, ...

in the interference patterns and the phase differences of the interference rays. For a double-ray interferometer  $\delta n = \frac{\lambda}{2\pi h} \frac{\delta I}{I_m}$ , for a multi-ray interferometer:  $\delta n = \frac{\lambda}{2hN_e} \frac{\delta I}{I_m}$ .  $I_m$  is the maximum intensity,  $h$  the

length of the ray in the gas to be examined,  $\delta I$  the intensity variation. With  $h = 10$  mm,  $\delta I/I_m = 5\%$  and  $R = 0.99$  ( $R$  - reflection coefficient of the mirrors)  $\delta n \leq 4 \cdot 10^{-7}$  for the double-ray instrument and  $\delta n \geq 4 \cdot 10^{-9}$  for the multi-ray instrument is obtained. The sensitivity thus differs by a factor  $B = \sqrt{R}/(1-R)$ . For  $R \rightarrow 1$ ,  $B \rightarrow \infty$ ; for  $R = 0.99$ ,  $B \approx 100$ . Maximum sensitivity is found to be obtained when  $I = 0.75 I_m$ . In practice,  $B$  will be affected by the external conditions and the errors of the interferometers.

ASSOCIATION: Kafedra optiki, Moskovskiy universitet (Department of Optics, Moscow University)

SUBMITTED: May 3, 1961 (initially)  
February 12, 1962 (after revision)

Card 2/2

KOROLEV, F.A.; KROMSKIY G.I.; SKOKOV, I.V.

Use of the phase method of multiwave interferometry for measuring  
low gas densities. Izv. vys. ucheb. zav.; fiz. no.5:61-63 '63.  
(MIRA 16:12)  
1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova.

KOROLEV, F.A.; KROMSKIY, G.I.; SKUROV, I.V.

Amplitude method of multiple-beam interferometry. Opt. i spektr. 14 no.3:  
416-418 Mr '63. (MIRA 16:4)  
(Interferometry)

AKIMOV, A.I.; KROMEKH, G.I.; SKOKOV, I.V.

Sensitivity of a multiple-wave interferometer. Prib. i tekhn.  
eksp. 9 no.5:172-174 S.O '64. (MIFI 17:12)

I. Fizicheskiy fakul'tet Moskovskogo gosudarstvennogo universiteta.

L 07066-67 EXT(1)  
ACC NR: AP6024633

SOURCE CODE: UR/0170/66/011/001/0003/0009

AUTHOR: Skokov, I. V.; Yemel'yanov, V. A.

ORG: none

TITLE: The use of multibeam interferometer for quantitative studies of axisymmetric inhomogeneities in rarefied gases

SOURCE: Inzhenerno-fizicheskiy zhurnal, v. 11, no. 1, 1966, 3-9

TOPIC TAGS: multibeam interferometer, gas flow, rarefied gas, gas density

ABSTRACT: A direct solution of the heat exchange problem during high speed (~5 km/sec) and high altitude (~120 km) flights is very difficult and thus it is of interest to develop diagnostic methods of densities in rarefied gases (Knudsen parameter  $\geq 0.01$ ). The author investigates rarefied gas density distributions around an axisymmetric model. The method of calculations of gaseous nonuniformities corresponding to slip flow are developed and an experimental device for the visualization of these events using multibeam interference is presented. The interferogram processing yields various parameters of the flow discussed. The density field in front of a sphere and the shock wave distribution for a disk are obtained in the

Card 1/2

UDC: 621.317.767

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ACC NR: AP6024633

case of a rarefied gas flow with  $M = 3.85$  and  $Re = 75$ . The experimental data are in fair agreement with the values obtained by the method of electron beam scattering (Ivanov, A. V. DAN SSSR, 161, 315, 1965). Orig. art. has: 17 formulas, 1 table, and 3 figures.

SUB CODE: 20/ SUBM DATE: 19Jan66/ ORIG REF: 007

Card 2/2 JC

ACC NR: AP7007727

SOURCE CODE: UR/0188/67/000/001/0110/0111

AUTHOR: Skokov, I. V.; Akimov, A. I.; Kromskiy, G. I.

ORG: MGU Department of Optics (MGU Kafedra optiki)

TITLE: Determination of shock wave profile by the interferometry method

SOURCE: Moscow. Universitet. Vestnik. Seriya III. Fizika, astronomiya, no. 1, 1967, 110-111

TOPIC TAGS: rarefied gas, gas density, gas dynamics, gas flow

ABSTRACT: The authors report the results of a study to determine using a multiple-wave interferometer the structure of the shock wave formed when rarefied gas moving at supersonic speed (Mach number  $\approx 4$ , Reynolds number = 50, stagnation temperature = 300°K) flows past a model (disk, diameter 10 mm). The investigated model was inserted between the mirrors of a Fabry-Perot etalon, which was illuminated by a collimated light beam from a point monochromatic light source, and the uniformly illuminated interference field was photographed. The negatives were processed using the photometric method taking axisymmetrical density distribution into account. The density distribution of the shock wave along the stagnation line in the vicinity of the forward critical point is shown in Fig. 1.

Card 1/3

UDC: 533.1:535.854

ACC NR: AP7007727

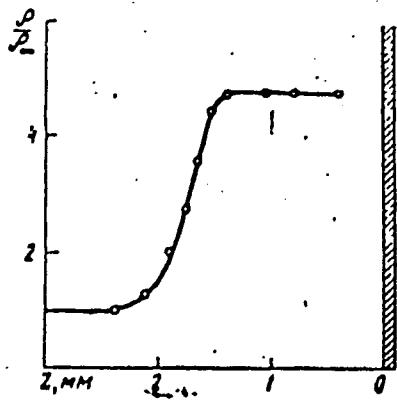


Fig. 1. Shock wave profile

$P/P_\infty$  is the ratio of current density value to the density of incoming flow;  
 $Z$  - is a coordinate along the flow axis.

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ACC NR: AP7007727

Under experimental conditions the thickness of the shock wave has a finite value ( $\sim 0.8$  mm) equal to approximately four lengths of the free path of particles of the incoming flow. The graph indicates clearly that there is a region with constant density. The ratio of densities in the incoming flow is in good agreement with Rankine's relationships (within limits of 10%). The relative value of the withdrawal of the shock wave is slightly higher than when the flow is continuous. It is noted in conclusion that the multiple-wave interferometry method permits determination of the density profile, the density fields near the model and in the free flow, the geometry of the shock wave, and other gas-dynamical parameters. Orig. art. has: 1 figure. [GS]

SUB CODE: 20 SUBM DATE: 16Jul66/ ORIG REF: 004/ ATD PRESS:5117

Card 3/3

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

SKOKOV, K.D.

Distortions in PS curves caused by electrolytic corrosion phenomena  
reported by OKS laboratories. Razved.i prom.geofiz.no.17:71-73 '57  
(MIRA 10:12)

(Seismometry)

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

SKOKOV, K.D.

Locating and eliminating leakages in logging cables. Razved. 1  
prom. geofiz. no.36:72-79 '60. (MIRA 13:12)  
(Oil well logging)

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

MAKAREVICH, V.N.; SKOKOV, E.D.

Some results of geothermal studies in deep wells of the Irtysh Valley. Razved. i issled. geofiz. no.189/7-102 163 (1981)

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

SKOKOV, L.D.; LOBIKOV, Yu.V.

Catch plate with floating cutters. Mashinostroyitel' no.11:21  
(MIRA 13:3)  
N '59.  
(Machine tools--Attachments)

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

L 05089-67 EWT(1)  
ACC NR: AP6013251

SOURCE CODE: UR/0413/66/000/008/0039/0039

20  
B

AUTHOR: Skokov, L. I.

ORG: none

TITLE: A waveguide-coaxial junction with a variable junction attenuation. Class  
21, No. 180664

SOURCE: Izobreteniya, promyshlennyye obraztay, tovarnyye znaki, no. 8, 1966, 39

TOPIC TAGS: waveguide coupler, coaxial cable, waveguide loss

ABSTRACT: This Author Certificate presents a waveguide-coaxial junction with a variable junction attenuation. The junction includes a section of a rectangular waveguide with two coaxial cables. The cables are located in one cross section of the rectangular waveguide on the wide walls and are connected to these walls by means of coupling probes. The coupling probes are extensions of the inner conductors of the cables. The design permits regulation of the junction attenuation within broad limits with matching from the waveguide. The coaxial cables are made so that they can be shifted along their axes. The cables are coupled by a

UDC: 621.372.833

Card 1/2

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ACC NR: AP6013251

mechanism which assures that when one of the cables in the waveguide is moved the other cable of the waveguide is advanced simultaneously. To obtain large values of the junction attenuations, the cables enter into the rectangular waveguide through terminal circular waveguides. In order to smooth the current distribution in the cables, the coupling areas mentioned above are fitted with matching cylindrical adapters fastened to their ends.

SUB CODE: 09/ SUBM DATE: 21Sep64

Card 2/2 LC

SKOKOV, N. (Stalino, USSR)

Michael Sarana guides a political study group. Pozh.dalo 5  
(MIRa 12:12)  
no.8:18-19 Ag '59.  
(Stalino--Communist education)

SKOKOWSKI, Tadeusz

Two sensitive strains of toxoplasmosis tests in various groups  
of the population. Sov. Med. 27 no.8:80-82 Ag '64.  
(MIR 1964)

1. Katedra infekcjonnykh bolezney s epidemiologiyey (zav.- prof.  
A.P. Petrov) Kuybyshevskogo meditsinskogo instituta.

10. *Urtica dioica* L. (Fig. 10)

and investigated and, in

Card 11

SKOKOVA, N. N.

"Ecology of the Gray Heron of the Rybinskiy Water Reservoir and Its Role in the Fish Economy of the Reservoir." Sub 5 Nov 51, Moscow City Pedagogical Inst imeni V. P. Potemkin.

Dissertations presented for science and engineering degrees in Moscow during 1951.

SO: Sum. No. 480, 9 May 55

SKOKOVA, M.N.

Results of banding Ibididae and Ardeidae in the U.S.S.R. Migr. zhiv.  
(MIRA 13:6)  
no.1:67-94 '59.

1. Astrakhanskiy zapovednik.  
(Herons) (Ibis) (Bird banding)

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

GERASIMOVA, T.D.; SKOKOVA, N.N.

Ornithogeographical characteristics of Aynov Islands.  
Ornitologiya no.2:91-98 '59. (MIRA 14:7)  
(Bol'shoy Aynov Island--Birds) (Malyy Aynov Island--Birds)

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

SKCKVA, N.N.

Ecology of the spoonbill and its role in the fishery management  
of the Volga Delta. Ornitologija no.2:262-270 '59. (MIRA 14:7)  
(Astrakhan Preserve--Spoonbills) (Birds--Food)

SKOKOVA, N.N.

Seasonal distribution and migrations of the large cormorant on  
the Caspian Sea. Migr. zhiv. no. 2:76-99 '60. (MIRA 13:12)

1. Astrakhanskiy gosudarstvennyy zapovednik.  
(Caspian sea region--Cormorants) (Birds--Migration)

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

SKOKOVA, N.N.

Feeding of the black-crowned night heron in the Volga Delta.  
Ornitologija no.3:396-404 '60. (MIRA 14:6)  
(Volga Delta--Herons)  
(Birds--Food)

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

SKOKOVA, N.N.

Food relationships of birds with colonial nesting habits in  
the coppices of the Volga Delta. Trudy Probl. i tek. sov.  
no.9:205-215 '60. (MIRA 13.9)

1. Astrakhanskiy gosudars tvennyy zapovednik.  
(Volga Delta--Birds--Food)

BIANKI, V.V., red.; KARPOVICH, V.N., red.; SKOKOVA, N.N., red.;  
KAS'YANOV, A.P., red.[deceased]; HELYAYEV, N.F., tekhn.  
red.

[Kandalaksha State Preserve] Kandalakshskii gosudarstvennyi zapovednik; nauchno-populiarnyi ocherk. Murmansk, Murmanskoe knizhnoe izd-vo, 1961. 87 p. (MIRA 16:6)

1. Kandalakshskiy gosudarstvennyy zapovednik.  
(Kandalaksha Preserve)

SKOKOVA, N.N.

Quantitative study of the feeding habits of ichthyophagous birds.  
Ornitologija no.4:288-296 '62. (MIA 16:4)  
(Astrakhan Preserve—Water birds) (Birds—Food)

SKOKOVA, N.N.

Puffins on the Aynovskiye Islands. Ornithologiya no. 5:7-12 '62.  
(MIRA 16:2)  
(Aynovskiye Islands—Puffins)

TARUSOV, B.N., otv. red.; SKOKOVA, N.N., red.

[Bioluminescence; a symposium of June 3-6, 1963;  
abstracts of reports] Bioluminestsentsiia; simpozium  
3-6 iiunia 1963 g.; tezisy dokladov. Moskva, 1963. 27 p.  
(MIRA 17:9)

1. Moskovskoye obshchestvo ispytateley prirody. Sektsiya  
biofiziki. 2. Moskovskiy gosudarstvennyy universitet (for  
Tarusov).

KONDRASHOVA, M.N.; Prinimali uchastiye: NIKOLAYEVA, L.V.; SKOKOVA, N.V.;  
SLEV, D.M.; TIMOFEEVA, L.M.

Effect of K-strophantin on phosphorylation and respiration of  
sarcosomes. Vop. med. Khim. 9 no. 3:273-279 My-Je '63.  
(MIRA 17:9)

1. Institut farmakologii i khimioterapii AMN SSSR i kafedra  
biokhimii zhivotnykh Moskovskogo gosudarstvennogo universiteta imeni  
Lomonosova.

CHONOWSKA-FUDOLIN H. Walka o zdrowie dziecka. The fight for children's health. stlaski Tygodnik Lekarski, Warszawa 1949, 4/29-30 (207-920) Tables 3

The war years were especially dangerous for the health and development of Polish children. The infection was three times and the mortality of babies twice that of before the war. To improve these conditions was the most important task of the Public Health Service. The number of dispensaries for pregnant women was continuously growing, and in 1946 reached 898 (from 254 in 1945). Their objects are the improving of the women's health and prenatal care. Every town and village is to have a midwife and a delivery room. A nursery for little children of preschool age will be established in every village. The mortality of babies has already remarkably decreased. It was 12.3 % in 1938, 26.9% in 1945, 13.3% in 1946 and only 9.7% in 1947. The medical service in the Primary Schools is extending. Vaccination against TB, diphtheria and other infectious diseases is carried out on a large scale. Moreover the following activities of New Poland in the fight for children's health are described: the dental service, the special surgical, orthopaedic, and laryngological dispensaries, the sanatoriums and preventoriums for TB, the holidays for healthy children in health resorts, milk supply for expectant and nursing mothers and for children up to the age of 14.

Makower - Wroclaw (IV, 7, 10)

So: Medical Microbiology and Hygiene, Section IV, Vol 3, No 1-6

Siodkowska - Rudolf, M

SIODKOWSKA-RUDOLF M.

Ochrona zdrowia matki i dziecka w planie sześciolatnim.  
Maternal-and child welfare in the Six Year Plan/ Pediat  
polska 24:5-6 May-June 50 p. 475-8.

1. NAI  
CIML Vol. 20, No. 2 Feb 1951

GORBUSHIN, M.F., inzh.; SKOKSHIN, L.V.

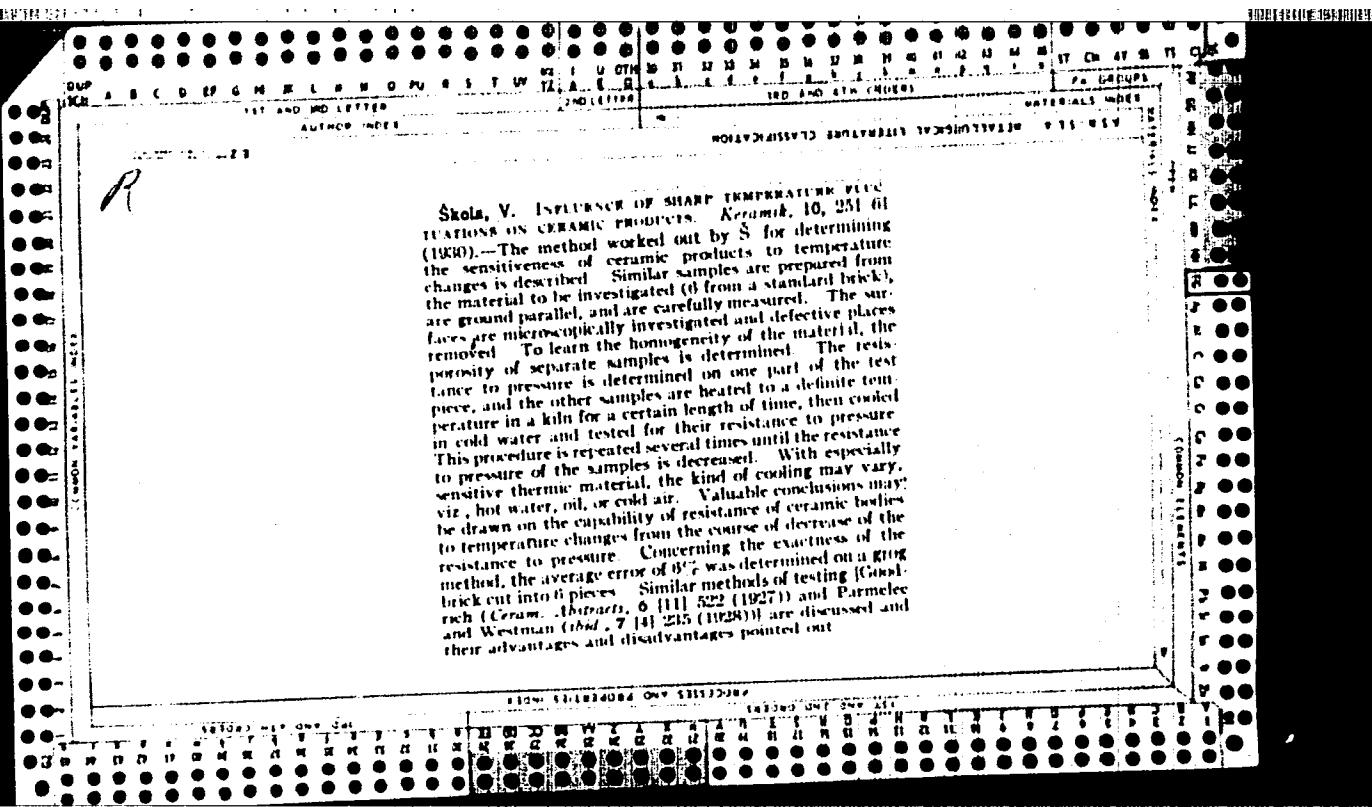
Chair conveyor for loading tires in freight cars. Mekh.i avtom.  
proizv. 16 no. 7:37-38 Jl '62. (MFA 15:6)  
(Conveying machinery)

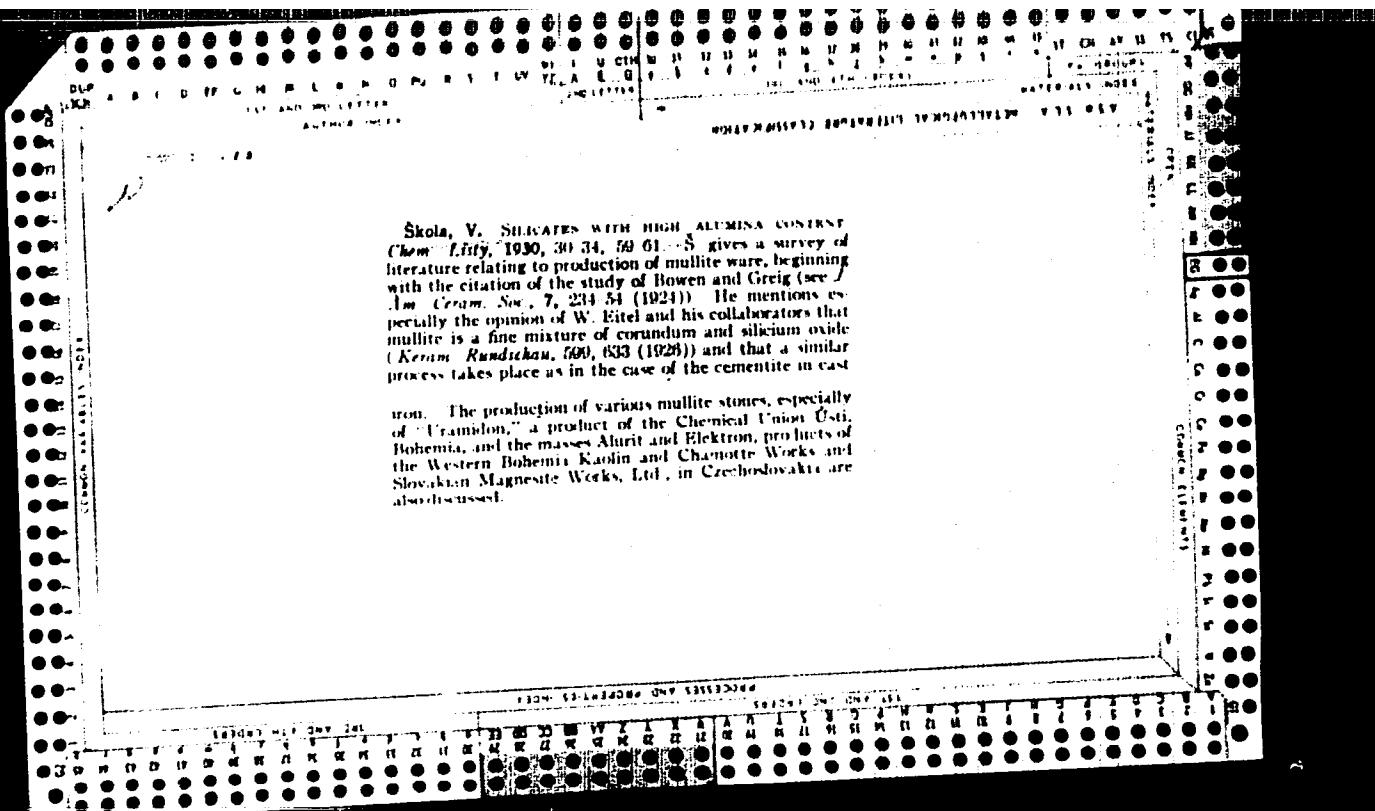
SKOLA, Jaroslav, prof., dr.

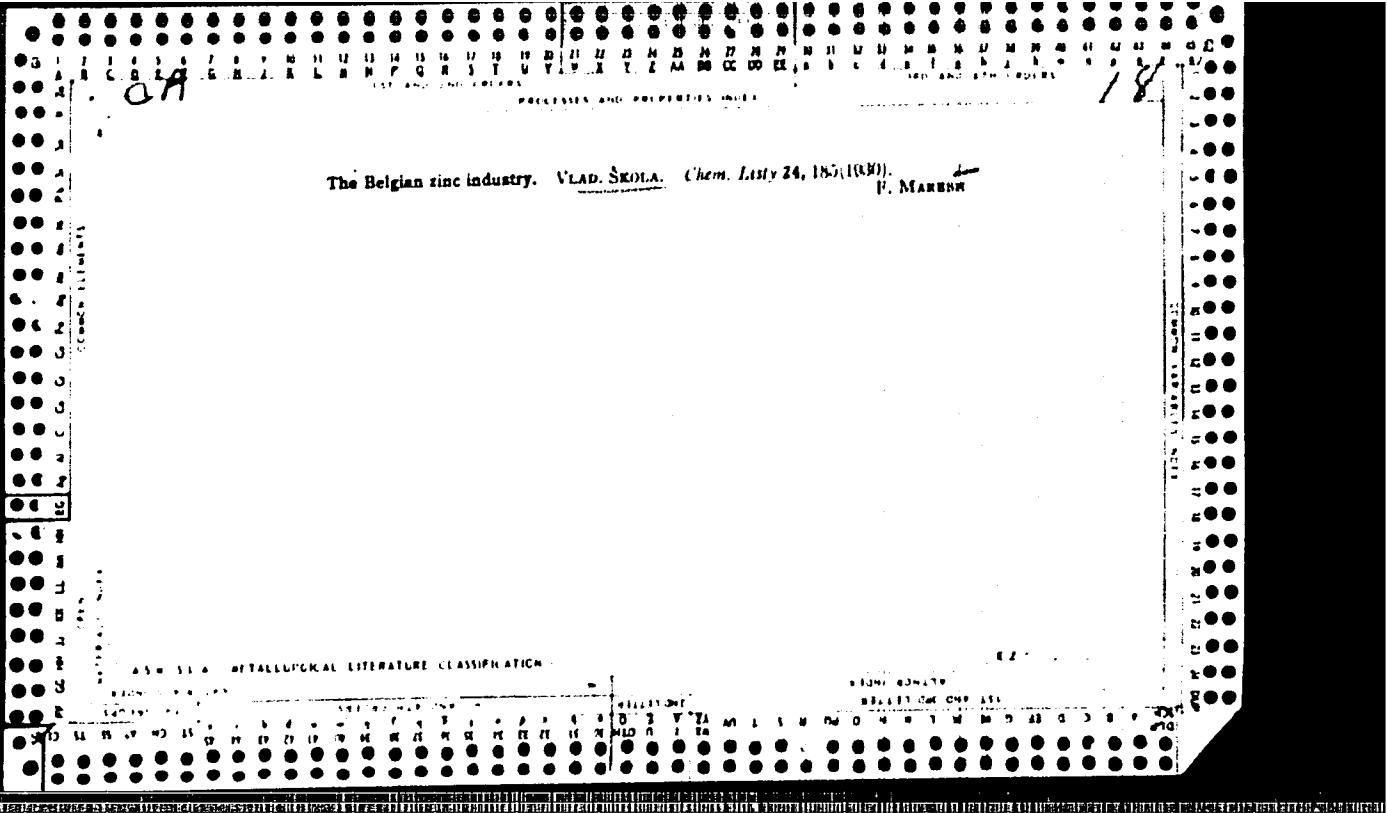
Problem of waste substances. Prum potravin 14 no.5:248 My '63.

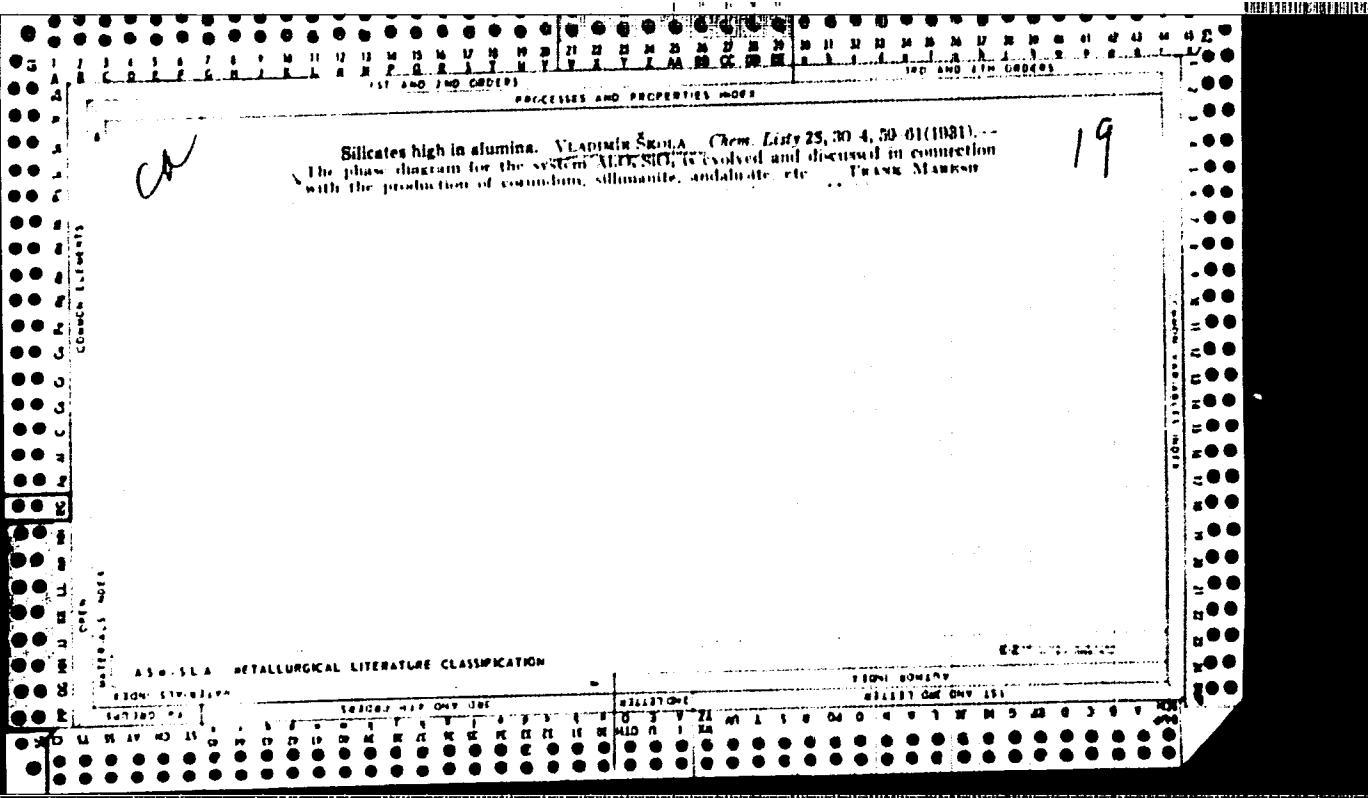
1. Ceskoslovenska akademie ved, Praha.

*[Signature]*  
Increasing the durability of refractory products. VLADIMIR ŠKOLA, Patent 1920, 508; Chem Abstract 9, 438.—By coating fireproof products with a cement consisting of oxides, their durability can be highly increased. One of the crucibles was coated on the inside with a 1-mm. thick Resistin coat. Both crucibles were then filled with the normal salt cake mixt. for mfg. sheet glass and heated in a test furnace for about 70 hrs. at a temp. of about 1350°. The crucible with a Resistin coat was not damaged at all, while the uncoated crucible was badly corroded. G.G.







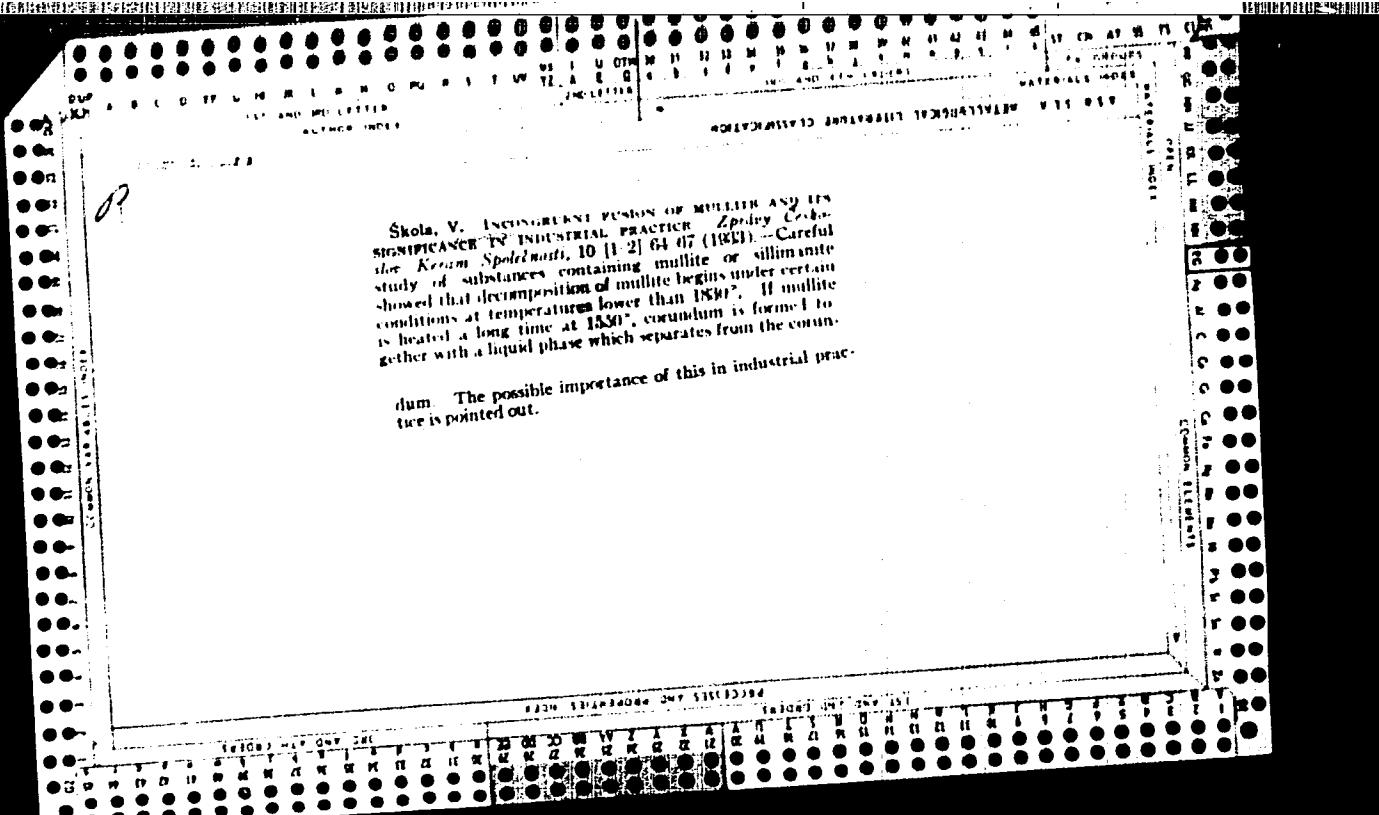


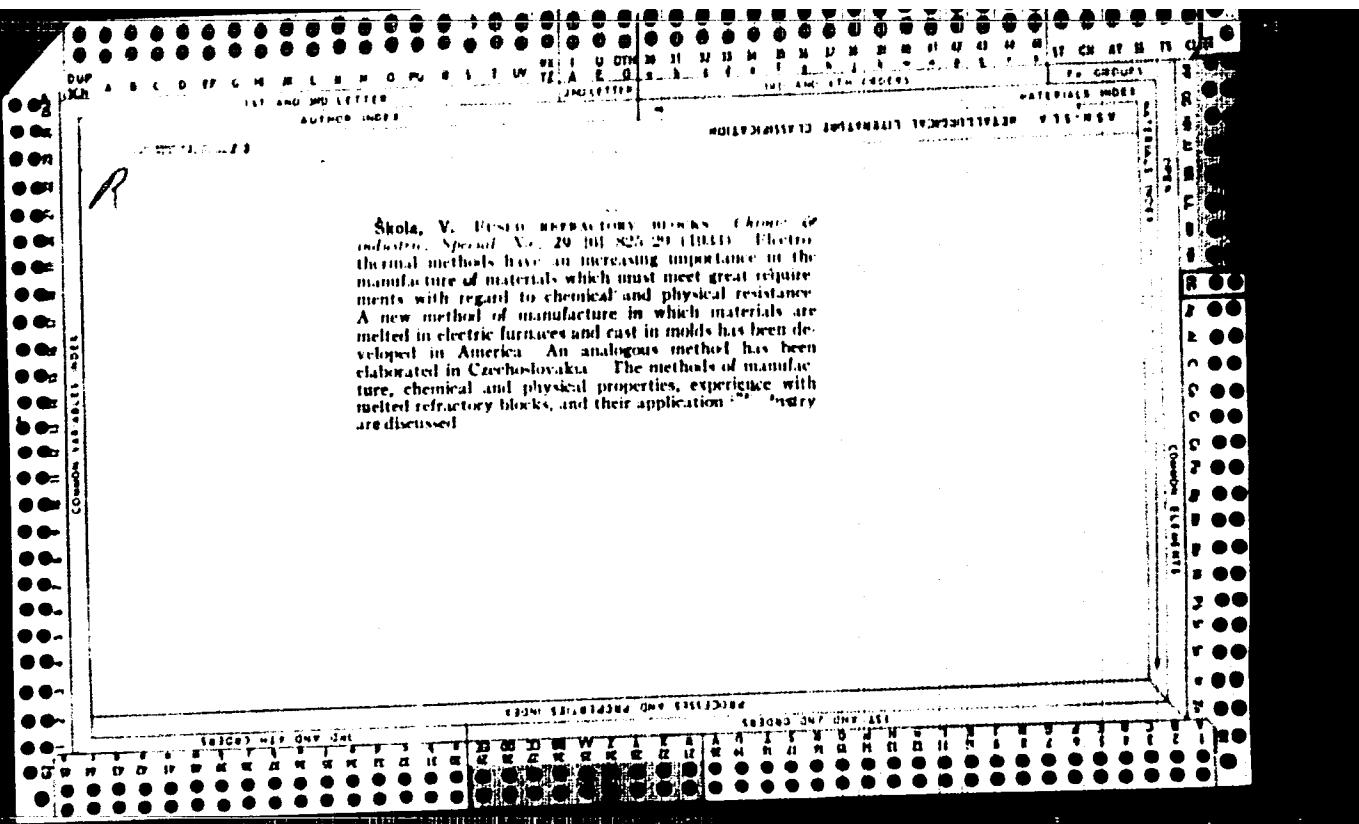
CA

28

The mannosidase fermentation of unextracted beet slices. NLAUDIA B. SCHAFFNER. *J. Am. Chem. Soc.*, 53, 165-71 (1931).—A complete analysis of fermented unextd. slices is given. The beet slices were pressed, and the juice after sterilization by heat was inoculated with a mannosidase enzyme. The fermented liquor was pptd. with EtOH and filtered through suspended filter bags. The clear filtrate was evapd., and the mannosidase crystallized. The mannosidase by osmosis was not successful. Com. trials have not been made. The literature is reviewed. FRANK MARSHALL

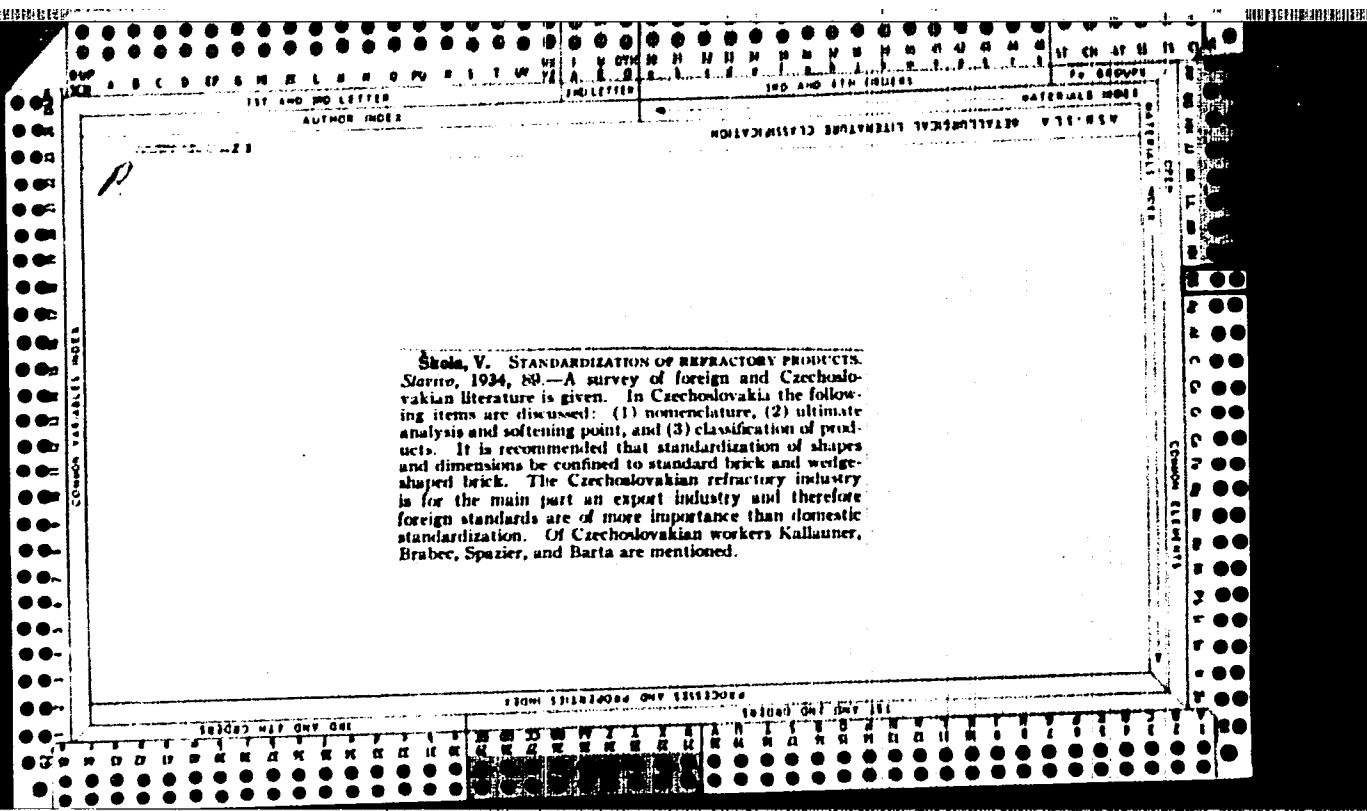
AMERICAN METAL PROCESS LITERATURE CLASSIFICATION

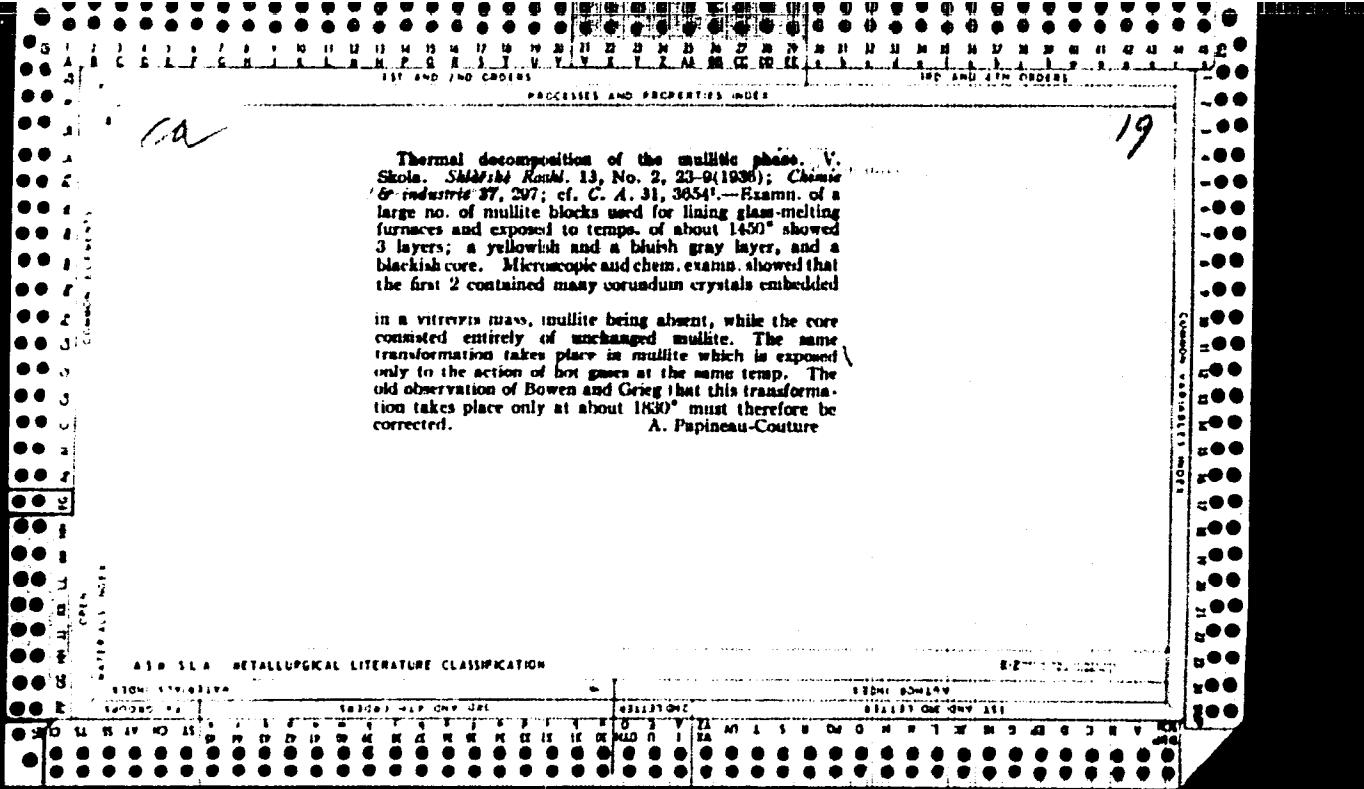




*✓* Fire-resistant materials in sugar establishments. Vladimír Skoda, Lotty Cukrová. No. 51, 405 500 (1983). Physico-chemical methods for testing refractory and enamel ware used in sugar mills, lime kilns and heating plants are surveyed. Frank Marsh

AS-15A METALLURGICAL LITERATURE CLASSIFICATION





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New methods for the production of refractory zirconium oxide ware. Vladimir Škoda. *Zprávy Českoslov. keram. společnosti* 12, 53-8 (1935); *Chem. Zentral.* 1936, II, 1904.—Bauxiteleyite is either fused in an elec. furnace with a high-quality clayey raw material (such as natural emery) and the melt poured into forms (corr. product, "Jargal") or the ore is finely ground, leached with  $H_2SO_4$ , the sulfate heated at a high temp., and  $ZrO_3$  obtained. Hot  $ZrC$  can also be treated with Cl and the resulting  $ZrCl_4$  converted into the oxide by heating at a high temp. M. G. Moore

M. G. Minge

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## ASB-SEA METALLURGICAL LITERATURE CLASSIFICATION

ANSWER

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

Physical and Chemical Properties, Testing, Etc.  
Disintegration of the Mullitic Phase at High Temperature.--V. Skola (Rospravy II tridy Ceske Akademie, 45, No. 17, 1935). According to Bowen and Greig Mullitic phases disintegrate at about 1,810° into corundum and a glass rich in silicic acid. The author found that disintegration may occur at a lower temperature, for example 1,450°, when alkalies are present. A mere 1% alkalies causes radical changes. If there is an excessive amount of silicic acid, mullite disintegrates with great difficulty even when alkalies are present. This is, for example, the case with mullite prepared by the thermal treatment of cyanite and sillimanite, and even more so with mullite isolated by hydrofluoric acid from burned sillimanite or with synthetic mullite.

R. B.

## ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION

EIRON SYNTHEZIVA

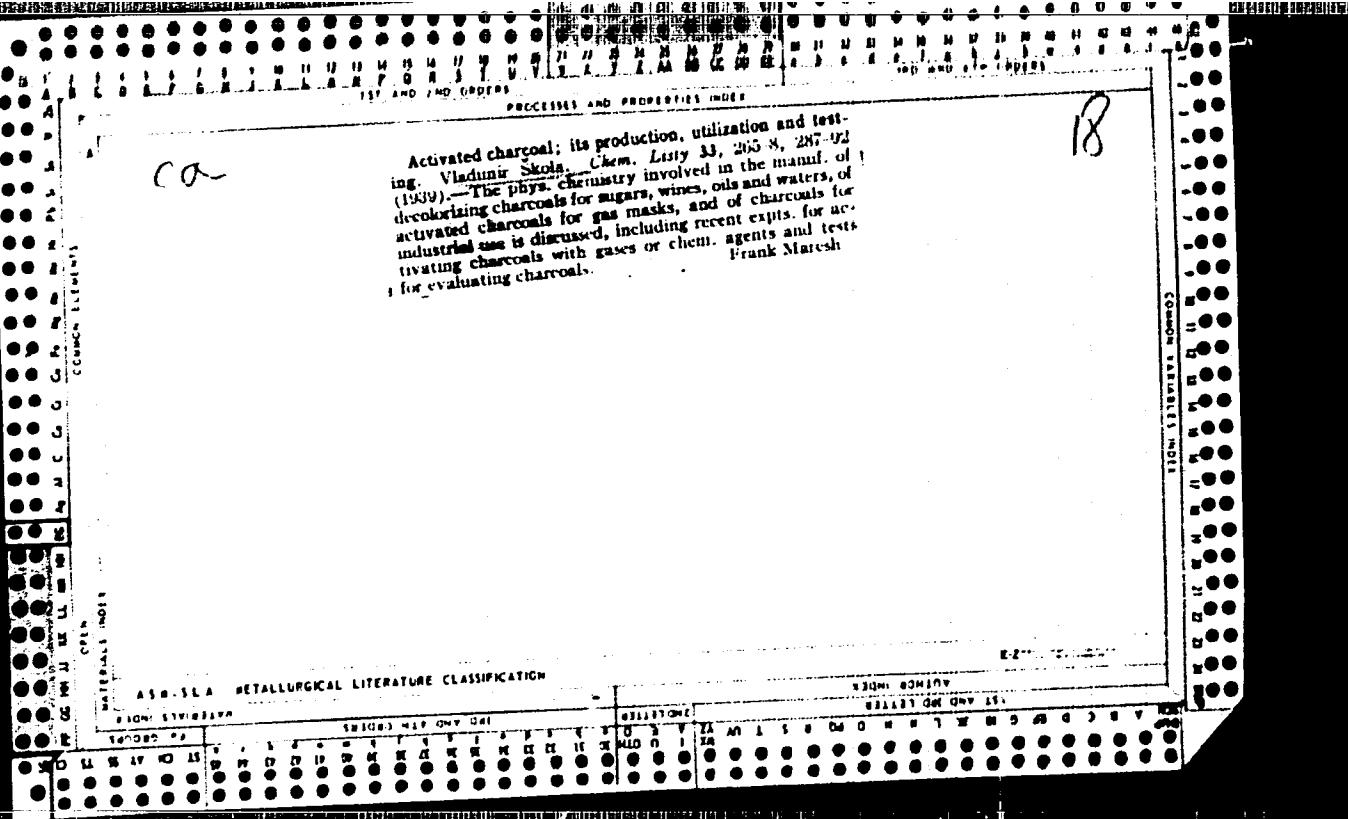
SANDOZ

193000 MEL ORE ORE

EIRON SYNTHEZIVA

SANDOZ

EIRON SYNTHEZIVA



SKOLA, V.

SKOLA, V. Use of nuclear reactors as a source of heat in chemical processes requiring high temperatures. Tr. from the English. p. 51.

Vol. 2, No. 2, Feb. 1957

NOVA TECHNICAL

TECHNOLOGY

Czechoslovakia

So. East European Accessions, Vol. 6, No. 5, May 1957

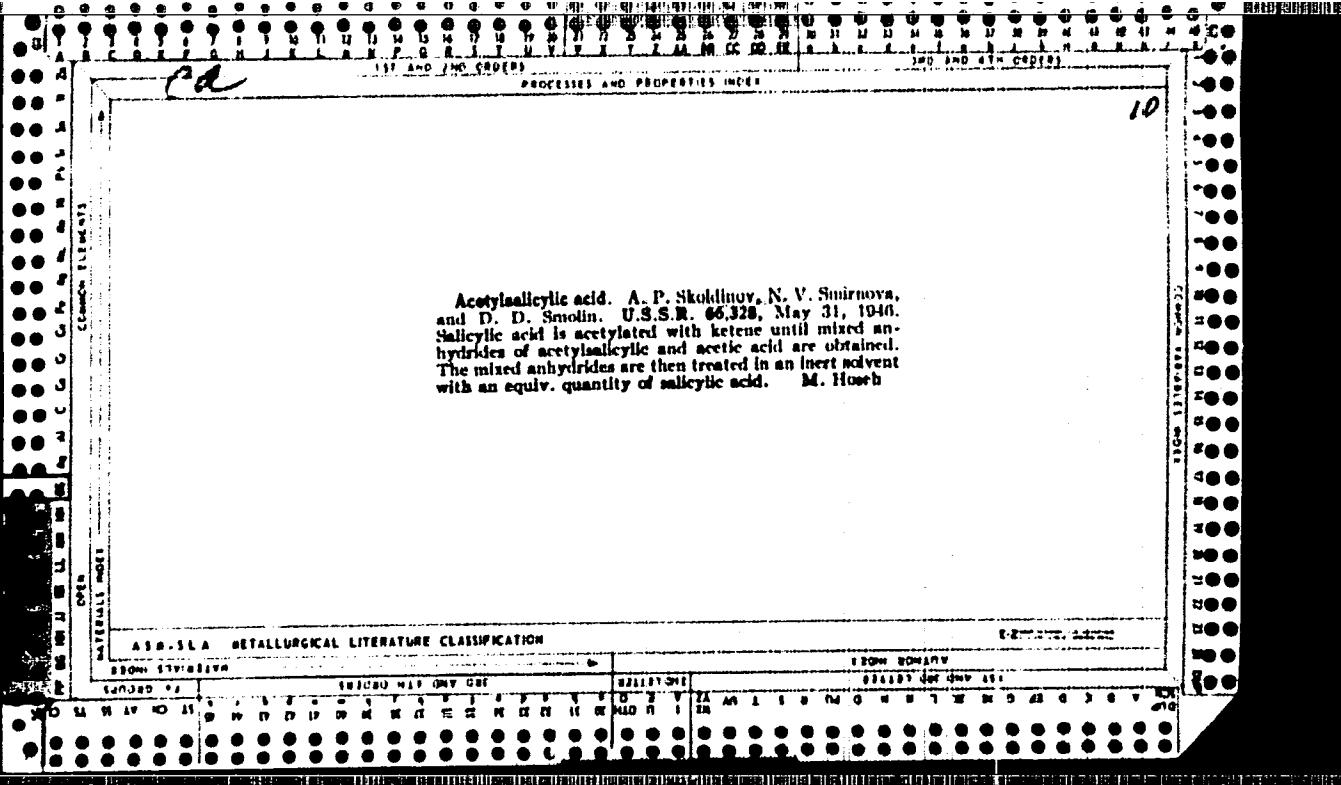
**Removing ferrous sulfate from copper sulfate.** A. P. Skoldinov, B. P. Kostev and N. M. Potashkin. Russ. 24,543, February 28, 1934.  $\text{FeSO}_4$  is oxidized by blowing air through the solution of  $\text{CuSO}_4$  in the presence of  $\text{CuCO}_3$ .

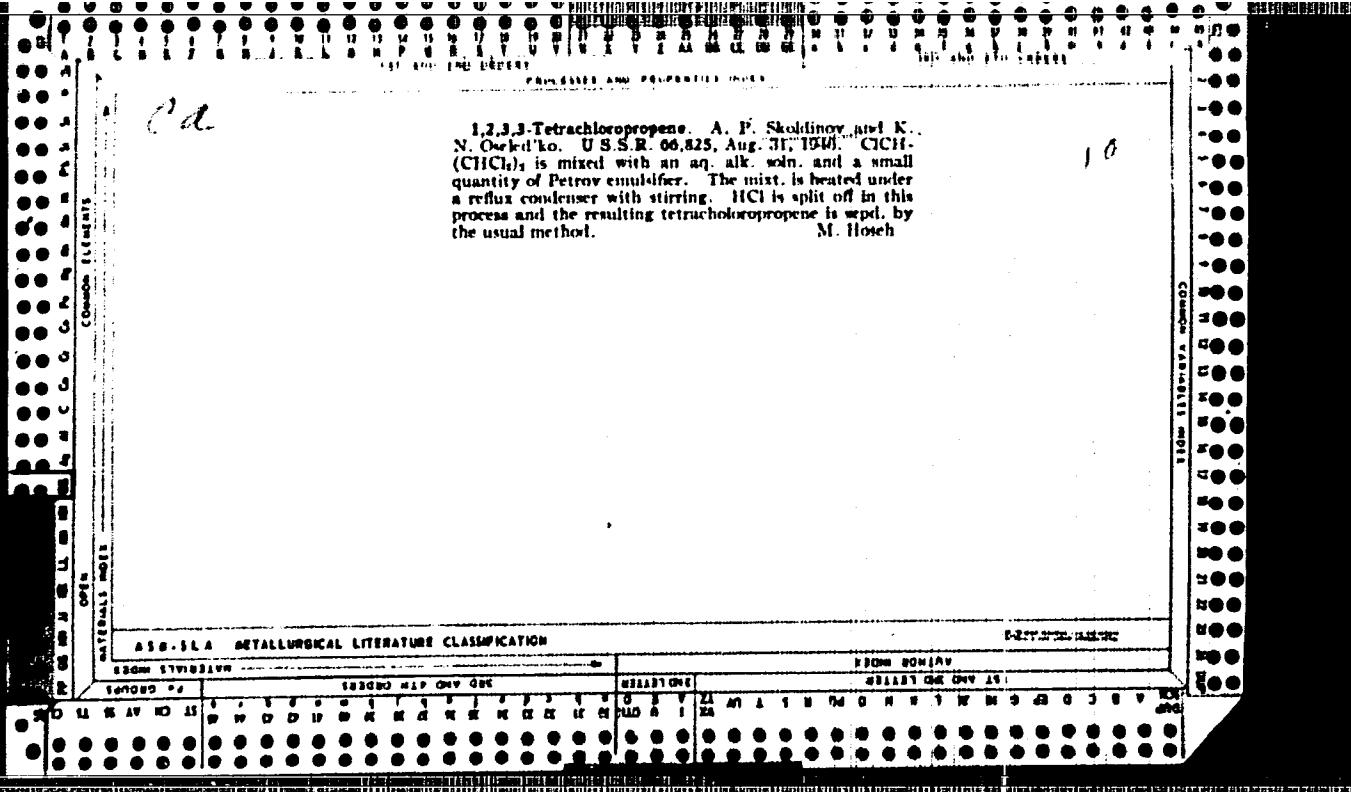
ASH-SEA METALLURGICAL LITERATURE CLASSIFICATION

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001551030002-4"

SKOLDINOV, A. P.

"Organometallic Compounds in the Friedel-Krafts Reaction," Zhur. Obshch. Khim., 12,  
No. 15, 7-8, 1942. Mbr., Lab. Agriculture, All-Union Inst. Exptl. Med. in. A. M.  
Gor'kiy, Moscow, -1941-.





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**Phenylurethans as raw materials for the preparation of sulfonamides and its derivatives.** V. A. Mikhalev and A. P. Skoldinov. *J. Applied Chem. (U.S.S.R.)*, 19, 1373-80 (1940).—Addition to C.A. 32, 811b. The following compounds were prep'd.: *p*-chlorophenylcarbamic acid, *Me* ester m. 110°; *Ei* ester m. 103-4°; *p*-sulfamylcarbamic acid, *Me* ester m. 229°, *Ei* ester m. 232°; *p*-(*p*-sulfamylphenylsulfamyl)carbamic acid, *Me* ester, *Ei* ester m. 253°; *p*-(2-pyridylsulfamyl)carbamic acid, *Me* ester m. 219-20°, *Ei* ester m. 210°; *p*-(4-methyl-2-thiazoylsulfamyl)carbamic acid, *Me* ester m. 243°, *Ei* ester m. 258°. R. Jones

ASA-SEA METALLURGICAL LITERATURE CLASSIFICATION

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APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001551030002-4"

Vibration spectra of acetylacetone derivatives, acetoacetic ester, and the hydrogen bond. D. N. Shigorin and A. P. Skoldinov (Karpov Phys.-Chem. Inst., Moscow). *Zhur. Fiz. Khim.*, 24, 955-97 (1950). — Neither  $\text{AcCH}(\text{R})\text{Ac}$  nor  $\text{AcCH}(\text{R})\text{CO}_2\text{Et}$  as liquid or in an inert solvent forms an intramol. H bond. The bands of the OH group of enolic isomers are characterized at 3350–3000  $\text{cm}^{-1}$ .

Paul W. Howerton

SKOLDINOV, A. P.

"Investigation in the Field of Acylation of Organic Compounds." Thesis for degree of Cand. Chemical Sci.  
Sub 16 Nov 19, Moscow Order of Lenin State U imeni M. V. Lomonosov.

Summary #2, 18 Dec 52, Dissertations Presented for Degrees in Science and Engineering in Moscow in 1949.  
From Vechernyaya Moskva, Jan-Dec 1949.

SKOLDINOV, A. P.

USSR/Chemistry - Production of Ketene

May 51

"Brief Communication: Large-scale Laboratory Apparatus for Production of Ketene,"  
D. D. Smolin, F. R. Kravtsov, A. P. Skoldinov, Sci Res Lab Exptl Chemotherapy,  
Min Public Health USSR

"Zhur Prik Khim" Vol XXIV, No 5, pp 547-551

Describes new large-scale lab ketene generator, made entirely of metal, yielding  
200-400 g/hr of ketene, depending on regime. It is based on principle of  
pyrolysis of acetone on open elec heating elements (spirals) placed in atm of  
acetone.

183T43

IVANOV, A.I.; SKOLDINOV, A.P.

Preparation of 1-(p-nitrophenyl)-2-aminoethanol and some derivatives.  
(MLRA 5:5)  
Zhur. Priklad. Khim., 25, 438-41 '52.  
(CA 47 no.14:6898 '53)

1. Ministry Health, Moscow.

SKOLDINOV, A. P.

USSR /Chemistry - Oxocompounds, Catalysts 21 MAY 52

"Acylation of Oxocompounds in the Presence of 'Acid' Agents," A. P. Skoldinov, K. A. Kochestkov, Cerr Mem, Acad Sci USSR, All-Union Sci Res Chem-Phar Inst imeni S. Ordzhonikidze

"Dok Ak Nauk SSSR" Vol LXXXIV, No 3, pp 527-530

This work reports the feasibility of progressive acylation of oxocompds in the presence of acid agents. Acid catalysts such as FeCl<sub>3</sub>, AlCl<sub>3</sub>, ZnCl<sub>2</sub>, SrCl<sub>4</sub>, P<sub>2</sub>O<sub>5</sub> could not be used, because of their violent reaction; but boron fluoride, which has a milder action and is more stable toward hydrolyzing

225T8

agents, led to the desired goal. It was shown that, in the progressive acetylation of oxocompds, all stages could be reached through the aid of acid agents, particularly boron fluoride and sulfuric acid. The C-acylation of oxocompds, by means of HF<sub>3</sub>, was achieved both immediately and by stages of the isomerization of O-acylate, thus leading to the assumption that even during the direct C-acylation process, the reaction takes place through an intermediate formation of O-acylate of enol, which is isomerized subsequently to C-acylate.

225T8

SKOLDINGOV, A. P.

232T10

USSR/Chemistry - Triacylnitrogen

1 Jun 52

"Preparation of Compounds of the Triacylnitrogen Type," N.V. Smirnova, A.P. Skoldinov, K.A. Kocheshkov, Cott Mem, Acad. of Sci SSSR, All-Union Sci Res Chem-Phar Inst imeni S. Ordzhonikidze

"Dok Ak Nauk SSSR" Vol 84, No 4, pp 737-740

Ketene reacts with amides in the presence of an inorg acid to form N-acetyl substituted amides. Further action of ketene on the diacetyl amide leads to the formation of compd having 3 acetyl groups on one nitrogen atom.

232T10

This compd is a representative of a new class of compds of the type  $N(COR)^3$  where R is an aliphatic radical. Nine compds of this series were prep and tabulated with their phys consts.

232T10

*Skoldinov, M.P.*

*p-Nitro- $\alpha$ -acetamido- $\beta$ -hydroxyacetophenone. A. P. Arzhangnik, M. I. Borukhova, V. A. Milkovets, O. I. Efimova, A. I. Skoldinov, D. D. Smidlin, and N. E. Smidina. U.S.S.R. 102,733, May 20, 1954. The title compound is obtained by the interaction of formaldehyde with  $p$ -nitro- $\alpha$ -acetamidoacetophenone in the presence of a condensation agent such as triethylamine.*  
*M. Weill*

*$\mu$ -Nitro- $\alpha$ -(dichloroacetylamino)acetophenone. A. P. Arzhangnik, A. I. Tsvetov, M. I. Borukhova, V. A. Milkovets, N. P. Skoldinov, D. D. Smidlin, and N. E. Smidina. U.S.S.R. 103,015, June 25, 1954. [Addn. to U.S.S.R. 102,733.] 1-Hydroxy-1-( $p$ -nitrophenyl)-2-aminoethane is treated with Cl<sub>2</sub>CHCO<sub>2</sub>Me and the resulting 1-hydroxy-1-( $p$ -nitrophenyl)-2-dichloroacetylaminoethane oxidized to  $\mu$ -nitro- $\alpha$ -(dichloroacetylamino)acetophenone.*  
*M. Weill*

SKOLDINOV, A. P.

Separation of a chlororaryl ketones from organic solvents.  
V. T. Klimko, N. K. Kochetkov, V. A. Mikulev, A. P. Skoldinov, and A. Ya. Khorlia; U.S.S.R. 103,767; Sept. 26, 1955. To solns. of chloro ketones RCOCH<sub>2</sub>Cl in org. solvents is added a tertiary amine and the ketones are sep'd as salts of quaternary NH<sub>3</sub> bases. M. Hoenig

Tetracycline. A. P. Arvidyanik and A. P. Skoldinov; U.S.S.R. 103,906; Sept. 25, 1955. Tetracycline is prepared by catalytic dehalogenation of 7-chlorotetracycline-HCl. The reaction is carried out over Pd catalyst in the presence of NH<sub>3</sub> or amines. M. Hoenig

SKOLDINOV, A. P.

Reaction of some derivatives of 9,9-dihydro-9H-  
propaanthracene. T. V. Protopova and A. P. Sko-  
ldinov. Zavod. Obrabotki Krem. 40, 1055-50 (1967).  
Celite was added at 60° to 1 hr; 13.8 g. CH<sub>2</sub>Cl<sub>2</sub> and  
1.8 g. HgO in 20 g. Celite after 3 hrs at 60°,  
yield 37% 1,4-dihydro-1-acetylphthalic anhydride  
99-8.5°, m. 242-6° [mp 103.5 (concentrated, m. 103-6°  
(decomp)]. Obtained from this ketone and CH<sub>2</sub>Cl<sub>2</sub>  
CH<sub>3</sub>NHCO<sub>2</sub> which is 70% yield treated with Na.

which contained 0.1 mole of Et<sub>2</sub>O, made blue, with 50% NaOH, and extd. with Et<sub>2</sub>O gave 81% 1-phenylpyrazole, b.p. 107-8°, also formed in 83% yield from I on treatment with NaOAc in aq. EtOH as above. II and NH<sub>4</sub>OH.HCl in 60% EtOH similarly gave 78% imidazole, isolated as CaCl<sub>2</sub> salt; I and NH<sub>4</sub>OH.HCl, with NaOAc, as above, gave 55% imidazole, also isolated as the CaCl<sub>2</sub> salt. To 3 g. I in 16 ml. EtOH was added 1.3 g. PPh<sub>3</sub>NH<sub>2</sub> and after refluxing 16 hr. the solid was collected on filter, washed with H<sub>2</sub>O and dried, giving 1.1 g. (31-33%) which was recrystallized from the 1/20 dilution of Et<sub>2</sub>O-H<sub>2</sub>O gave 1.04-1.05 g. II (44-47%).

aq. NaOH and heat. with hot C<sub>6</sub>H<sub>6</sub>. 70% 2-aminopropylidene, m. 135-6°. II similarly gave 71% yield.

KUL DE NOV, H. E.

*V<sub>B</sub>-Acetylacrolein* A. P. Skoldinsky and T. V. Tikhonova. U.S.S.R. 106,370, July 25, 1957. An equ. soln. of Na malondialdehyde or a suspension of its anhyd. with an inert solvent is treated with a carboxylic acid chloride and the reaction product is sepd. by known means. M. H. S.

451

AUTHORS:

S A T L D , N C Y , 1 ) 1  
Protopopova, T. V., and Skoldinov, A. P.

TITLE:

1,1,3,3-Tetraalkoxy Propanes (1,1,3,3-Tetraalkoxipropany)

PERIODICAL:

Zhurnal Obshchey Khimii, 1957, Vol. 27, No. 1, pp. 57-62 (U.S.S.R.)

ABSTRACT:

The properties of 1,1,3,3-tetraalkoxy propanes obtained during the reaction of vinyl ethers with orthoformic acid esters in the presence of a Friedel-Crafts type catalyst, are analyzed. This reaction is similar to the well-known reaction of vinyl ethers with acetals leading to the formation of alkoxy acetals or products of further addition of one or several vinyl ether molecules. The addition of the orthoformic acid ester to the vinyl ether results not only in the formation of 1,1,3,3-tetraalkoxy propane but the latter in turn can attach itself by its double bond, forming 1,1,3,5,5-pentaalkoxy pentane and products of further reaction with various molar ratios of the orthoformic acid ester and vinyl ether and the yield of the mono-additive in this case depends less upon the surplus ester used than in the case of addition of acetals. This is explained by the fact that the orthoester function during the reaction of boron trifluoride becomes ionized much easier than the acetal function. The relatively greater

Card 1/2

KLIMKO, V.T.; KHLORLIN, A.Ya.; MIKHALEV, V.A.; SKOLDINOV, A.P.; KOCHETKOV, N.K.

$\beta$ -aminovinyl ketones. Part 7: Reaction of  $\beta$ -chlorovinyl ketones  
with tertiary amines. Zhur. ob. khim. 27 no.1:62-65 Ja '57.

(MLRA 10:6)

I. Institut farmakologii i khimioterapii Akademii meditsinskikh nauk  
SSSR.

(Vinyl compounds) (Ketones) (Amines)

SAC DENNIS

79-2-20/58

AUTHORS:

Klimko, V. T.; Mikhalev, V. A.; Skoldinov, A. P.

TITLE:

Investigation of Derivatives of beta-Dicarbonyl Compounds. Part 1.  
Synthesis of beta-Chlorovinylketones. (Issledovaniya v oblasti  
proizvodnykh beta-dikarbonilnykh soyedineniy. I. Sintez beta-khlorvinyl-  
ketonov)

PERIODICAL:

Zhurnal Obshchey Khimii, 1957, vol 27, No 2, pp. 370-374 (U.S.S.R.)

ABSTRACT:

The authors improved a method of obtaining beta-chlorovinyl ketones by the condensation of carbonyl chlorides with vinyl chloride and enlarged the field of application of this synthesis. The intermediate products of this condensation were identified as beta-beta-dichloroethyl ketones. It was established that the intermediately forming aryl-beta-beta-dichloroethyl ketones have a much higher stability than their aliphatic analogues. An attempt to obtain beta, beta-dichloroethylaryl ketone by the reaction of p-methoxybenzoyl chloride with vinyl chloride in the presence of anhydrous aluminum chloride was unsuccessful because the methoxyl group was also involved in the reaction. By using anhydrous aluminum chloride in nitromethane in the role of the condensation agent, it was possible to obtain a 60% yield

Card 1/2

79-2-20/58

Investigation of Derivatives of beta-Dicarbonyl Compounds. Part 1.

of p-methoxyphenyl-beta, beta-dichloroethyl ketone. The obtained aryl-beta, beta-dichloro ketones were quite stable in storage and slowly separated the hydrogen chloride during the boiling with aqueous bicarbonate or sodium carbonate solutions. The conversion of beta-beta-dichloroethyl ketones into homologous beta-chlorovinyl ketones was realized easily during the reaction of the former with trialkylamines (e. g., triethylamine).

There are 13 references, of which 7 are Slavic.

ASSOCIATION: USSR Academy of Medical Sciences, Institute of Pharmacology and Chemotherapy

PRESENTED BY:

SUBMITTED: January 9, 1956

AVAILABLE: Library of Congress

Card 2/2

SKOLDINOV, A.P.

✓ Synthesis with 1,1,2,2-tetraethoxyethane. Preparation of the ring of the compound containing the Protoporphyrin IX ring system. - Urea (0.1 g.), 110 ml. EtOH, and 22 g.  $\text{CH}_3\text{C}(\text{Cl})(\text{OBu})_2$  (1) treated with 20 ml. concn. HCl and the mixture kept overnight and cooled. *in vacuo* yielded 72% 2-hydroxyprotoporphyrin IX.

ARENDARUK, A.P.; BUDOVSKIY, E.I.; GOTTIKH, B.P.; KARPEYSKIY, M.Ya.  
KUDRYASHOV, L.I.; SKOLDINOV, A.P.; SMIRNOVA, N.V.; KHORLIN, A.Ya.  
KOCHETKOV, N.K.

Dihydrosarcomycin and related compounds. Part.1. Zhur.ob.khim.  
27 no.5:1312-1318 My '57. (MLRA 10:8)

1.Institut farmakologii i khimioterapii Akademii meditsinskikh  
nauk SSSR.  
(Antibiotics)

PROTOPOPOV, T.V.; SKOLDINOV, A.P.

Production and some transformations of 1-alkoxy-1,3,3-trihalide  
propanes. Khim. nauka i prom. 3 no.4:536 '58. (MIRA 11:10)

1. Institut farmakologii i khimioterapii AMN SSSR.  
(Propane)

SKOLDINOV A.P.

ARENDARUK, A.P.; SKOLDINOV, A.P.

Production of tetracycline. Med.prom. 12 no.4:14-17 Ap '58.  
(MIRA 11:5)

1. Institut farmakologii i khimioterapii Akademii meditsinskikh  
nauk SSSR.  
(TETRACYCLINE)

SMIRNOVA, N.V., ARIENDARUK, A.P., SNOLIN, D.D., SKOLDINOV, A.P.

Esters of N-(arylalkyl)-4-phenylisonipeptic acid. Med.prom.12 no.7  
(MIRA 11:8)  
31-35 J1 '58

1. Institut farmakologii i khimioterapii AMN SSSR.  
(NIPPECOTIC ACID)

SKOLDINOV, A. P.

79-1-54/63

AUTHORS: Protopopova, T. V., Skoldinov, A. P.  
TITLE:  $\beta$ -Acetoxyacroleins ( $\beta$ -Atsiloksiakroleiny)  
PERIODICAL: Zhurnal Obshchey Khimii, 1950, Vol.20, Nr 1, pr.240-243(USSR)

ABSTRACT:  $\beta$ -Acylolymuroleins of the general formula (AcOCCH = CH-CH=O) (I) had hitherto not been described. They contain a reactive system of  $\alpha$ , $\beta$ -unsaturated aldehyde and may serve as initial products for interesting syntheses. The authors obtained them by conversion of the sodium salt of malonic dialdehyde (II) with chlorine anhydrides of acids. With the use of chlorine anhydrides of acetic, propionic, n-butyric or benzoic acid the method A gave good results. It consisted of the conversion of the chlorine anhydride with a suspension consisting of the anhydrous sodium salt of malonic dialdehyde in an inert solvent. The initial salt (II) was obtained by saponification of 1,1,3,3-tetraoxyp propane in the presence of hydrochloric acid and by subsequent neutralization with caustic soda. The reactions with the chlorine anhydride of p-nitrobenzoic acid take place better according to method B' which

Card 1/5

70-1051/63

$\beta$ -Acyloxyacroleins

consists of the influence of chlorine anhydride upon the aqueous solution of the salt (II). The  $\beta$ -acyloxyacroleins which contain the residue of an aliphatic acid are liquids of a pungent specific odor which are distillable in a vacuum and which show molecular exaltation. But the compounds with an aromatic acyl residue are crystalline bodies. All  $\beta$ -acylacrleins are soluble in the usual organic solvents with the exception of petroleum ether, insoluble in water and aqueous soda solutions, but they are easily saponified by the latter under formation of malonic dialdehyde. They show the typical reactions upon the aldehyde group and easily condense with salts of guanidine under formation of 2-aminopurimidine. All described properties prove the structure (I) and are incompatible with the structure of acylmalonic dialdehydes  $AC\ CH(CH=O)_2$  which might be expected when the acylation process would be accompanied by a transfer of the reaction center. (See tables 1 and 2). There are 2 tables, and 2 references; all of which are Slavic.

Card 2/3

$\beta$ -Acyloxyacroleins

79-1-51/63

ASSOCIATION: Institute for Pharmacology and Chemotherapy of the Academy  
of Medical Sciences, USSR  
(Institut farmakologii i khimoterapii Akademii meditsinskikh  
nauk SSSR)

SUBMITTED: December 28, 1956

AVAILABLE: Library of Congress

Card 3/3

1. Chemistry 2. Anhydrides 3. Dialdehydes 4. Chemical reactions-  
Theory

PROTOPOPOVA, T.V.; SKOLDINOV, A.P.

Acetals of  $\beta,\beta$ -dihalogen substituted propionaldehyde. Zhur. ob. khim.  
28 no.9:2428-2431 S '58. (MIRA 11:11)

1. Institut farmakologii i khimioterapii AMN SSSR.  
(Propionaldehyde)

AUTHORS: Protopopova, T. V., Skoldinov, A. P. SOV/79-26-10-36/60

TITLE:  $\beta\beta$ -Dihalogen Propionic Aldehydes ( $\beta,\beta$ -Digalcidopropionovye  
al'degidy)

PERIODICAL: Zhurnal obshchey khimii, 1958, Vol 28, Nr 10,  
pp 2805-2808 (USSR)

ABSTRACT: These aldehydes  $\text{CHX}_2\text{CH}_2\text{CHO}$ , where X=Cl or Br (Compound I). have not hitherto been obtained in free state (Ref 1). Also the authors did not succeed in synthesizing them by the hydrolysis of the corresponding acetals  $\text{CHX}_2\text{CH}_2\text{CH}(\text{OR})_2$ , where R=alkyl (III)(Ref 2). As previously reported (Ref 2) the acetals (III) are easily transformed into the corresponding 1,1,3-trihalogen-3-alkoxy propane  $\text{CHX}_2\text{CH}_2\text{CHX}' (\text{OR})(\text{IV})$  by the action of phosphorus pentachloride or thionyl chloride. When these reactive compounds are shaken with water homogeneous aqueous solutions of the aldehydes (I) are obtained, wherefrom the latter are separated; they can then be further distilled in the nitrogen current in a vacuum. The aldehydes (I) may be synthesized

Card 1/3

$\beta,\beta$ -Dihalogen Propionic Aldehydes

SOV/79-28-1e-36/6c

in the same way from the compounds (II), although in smaller yields and a longer period of time being necessary. The aldehydes  $\text{CH}_2\text{CH}_2\text{CHO}$ (I) are liquids of a strong specific smell; they decompose already at room temperature under the formation of hydrogen halide and the formation of solid polymers; the cleavage is slowed down by hydroquinone. The analysis of the  $\beta,\beta$ -dibromopropionic aldehyde, which is less stable than the chlorine analog was, therefore, difficult to carry out. The compounds (I) show the characteristic reactions to the aldehyde group and form crystalline unstable derivatives that can only be recrystallized in the absence of acids. In the oxidation of (I)(X=Cl) with concentrated nitric acid the  $\beta,\beta$ -dichloro propionic acid was obtained. There are 3 references, 1 of which is Soviet.

ASSOCIATION: Institut farmakologii i khimioterapii Akademii meditsinskikh nauk SSSR (Institute of Pharmacology and Chemotherapy of the Academy of Medical Sciences USSR)

Card 2/3

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4

$\beta,\beta$ -Dihalogen Propionic Aldehydes

SOV/70-26-10-31/60

SUBMITTED: September 30, 1957

Card 3/3

APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001551030002-4"

Mikhailov, V.A.; Dorokhova, M.I.; Smolina, N.Ye.; Zhelezovtseva, A.M.;  
Tikhonova, O.Ya.; Skoldinov, A.P.; Arendaruk, A.P.; Smolin, D.D.;  
Golovkina, T.V.; Slonova, L.A.

Styrene as an initial product for synthomycetin and levomycetin  
production. Part 2: Synthesis of p-nitroacetophenone and  
p-nitro- $\alpha$ -bromacetophenone. Antibiotiki 4 no.4:21-24 Jl-Ag  
'59.  
(MIRA 12:11)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevicheskiy  
institut imeni S.Ordzhonikidze (for Mikhailov, Dorokhova, Smolina,  
Zhelezovtseva, Tikhonova). 2. Institut farmakologii i khimio-  
terapii AMN SSSR (for Skoldinov, Arendaruk, Smolin, Golovkina,  
Slonova).

(CHLORAMPHENICOL chem)  
(KETONES chem)

17294  
207/63-4-6-23/37

AUTHORS: Protopopova, T. V., Klimko, V. T., Skoldinov, A. P.

TITLE: Brief Communications. Substituted Malondialdehydes and Some of Their Reactions

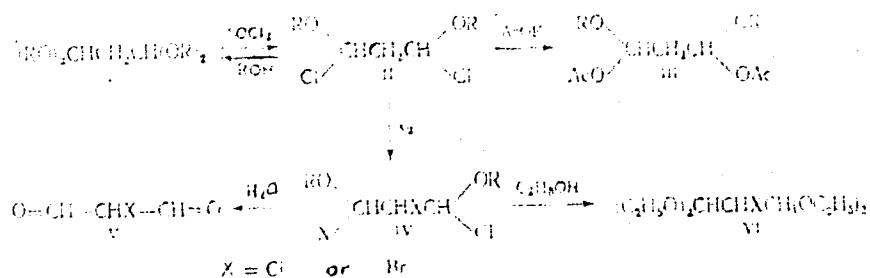
PERIODICAL: Khimicheskaya nauka i promyshlennost', 1959, Vol 4, No 6, pp 305-306 (USSR)

ABSTRACT: The preparation and halogenation of 1,3-dichloro-1,3-dialkoxypropanes (II) was investigated. Di- $\alpha$ -chloroethers (II) are easily formed from acetals (I) and thionyl chloride or phosphorus pentachloride. II with alcohols are converted into I and by action of  $\text{CH}_3\text{COOK}$ , diacetohydroxy derivatives (III) are formed.

Card 1/5

Brief Communications  
Sulfoxidized Malondialdehydes and  
Some of Their Reactions

77264  
367/63-4-6-36/37



The yields and properties of acetals of the homologs  
of malondialdehyde are given in Table 2.

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Brief Communications. Substituted  
Malondialdehydes and Some of Their  
Reactions

77294  
SOV/63-4-6-26/37

R	R'	YIELD (%)	BOILING POINT (°C)	$n_D^{20}$	$\delta_4^{20}$	MELTING POINT OF DIANYL HYDROCHLORIDE (PC)
CH <sub>3</sub>	CH <sub>3</sub>	53.4	87-87.5/5 mm	1.4151	0.9158	223-224
CH <sub>3</sub>	CH <sub>3</sub>	39.8	62-63/5 mm	1.4090	0.9652	223-224
C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	64.0	82-83/3 mm	1.4179	0.9097	206-207*
iso-C <sub>3</sub> H <sub>7</sub>	C <sub>2</sub> H <sub>5</sub>	68.0	80-81.5/2 mm	1.4220	0.9107	167-171
C <sub>6</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	43.9	125-127/1.5-2 mm	1.4715	0.9826	143-144

\*LITERATURE SOURCES: MP 206-207

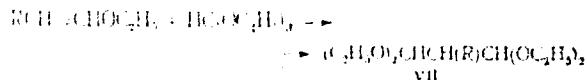
Table 2. Acetals of the homologs of malondialdehyde  
 $(RO^1)_2CHCHRCH(OR^1)_2$ .

Card 3/5

Baker Communications. Substituted  
Malondialdehydes and Some of Their  
Reactions

77294  
30V/63-4-6-28/37

The complete acetals of the homologs of malondialdehyde (VII) were obtained by condensation of alkenyl ethyl ether with orthoformic ester in the presence of acid catalyst.

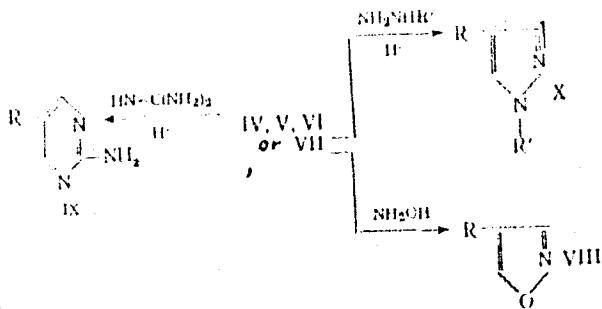


Substituted malondialdehydes (V), their derivatives (IV) and (VI), also acetal (VII) with hydroxylamine, guanidine, or with hydrazines, form 4-substituted isoxazoles (VIII), corresponding 5-substituted 2-aminopirimidines (IX), or 4-substituted pyrazoles (X).

Card 4/5

Brief Communications. Substituted  
Methylenepyrazoles and Some of Their  
Reactions

77294  
307/25-4-6-23/37



The yields and boiling points of 4-substituted pyrazoles are given in Table 5. There are 5 tables; and 3 references, 1 Soviet, 1 Japanese, 1 U.S. The U.S. reference is: C. Swallen, E. Board, J. Am. Chem. Soc., 52, 654 (1930).

Card 5/5    Ind. Pharmacology, Chemotherapy, AMS USSR

AUTHORS:

Solov'yev, V. M., Arendaruk, A. P.,  
Skoldinov, A. P.

SOV/79-29-2-58/71

TITLE:

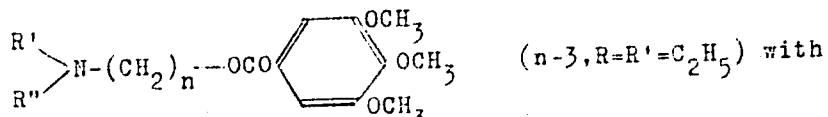
Dialkylaminoalkyl Ester of 3,4,5-Trimethoxy Benzoic Acid  
(Dialkilaminoalkilovye estiry 3,4,5-trimetoksibenzoynoy kisloty)

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 2, pp 631-635 (USSR)

ABSTRACT:

Reserpine (I), a high-active hypotensive tranquilizing agent, is the most important alkaloid extracted from the plant Rauwolfa and is generally used in medicine. Recent reports stated that there is a great number of simpler compounds, possessing only a part of the structural elements found in reserpine but having the same activity. Thus, according to data contained in publications, compound (II) shows one third of the reserpine activity and it is assumed that the activity is caused by the similarity of the structure of (II).



the structure of a molecular particle of reserpine (Ref 1).

Card 1/3

Dialkylaminoalkyl Ester of 3,4,5-Trimethoxy Benzoic Acid SOV/79-29-2-58/71

If this assumption is right, it must be possible to synthesize relatively simply structured compounds possessing the above mentioned properties of reserpine. Therefore the authors synthesized 12 dialkylaminoalkyl esters of 3,4,5-trimethoxy benzoic acid (II) differing as to the length of the carbon chain between the atoms of oxygen and nitrogen, as well as to the substituents at the nitrogen atom (Table). All esters apart from (II)(n=3, R=R'=C<sub>2</sub>H<sub>5</sub>) were synthesized by the reaction of equimolecular quantities of chloric anhydride of 3,4,5-trimethoxy benzoic acid (Ref 2) with the corresponding amino alcohol in benzene medium. The bases, which were easily obtained in pure state, were characterized as chloro hydrates and iodine methylates. 2-(N-hexamethylene imino)-ethanol-1, which is not described in publications, was obtained, like other amino alcohols, with n = 2, by the reaction of hexamethylene with ethylene bromo hydrin. Among the compounds synthesized, only a part develops a limited activity. It was not possible to find a relation between reserpine and the structural similarity and the activity of the preparations obtained. Pharmacological

Card 2/3

Dialkylaminoalkyl Ester of 3,4,5-Trimethoxy Benzoic Acid SOV/79-29-2-58/71

investigations have yet to been undertaken. There are 1 table and 9 references, 2 of which are Soviet.

ASSOCIATION: Institut farmakologii i khimioterapii Akademii meditsinskikh nauk SSSR (Institute of Pharmacology and Chemotherapy of the Academy of Medical Sciences, USSR)

SUBMITTED: December 25, 1957

Card 3/3

5 (3)

AUTHORS: Protopopova, T. V., Skoldinov, A. P. SOV/79-29-3-45/61

TITLE:  $\beta$ -Halogen Acroleins ( $\beta$ -Galoidoakroleiny)

PERIODICAL: Zhurnal obshchey khimii, 1959, Vol 29, Nr 3, pp 963-967 (USSR)

ABSTRACT: Proceeding from the synthesis of  $\beta,\beta$ -dihalogen propionaldehydes (Refs 1,2) the authors attempted in the work under review to transform these aldehydes (I) into the corresponding  $\beta$ -halogen acroleins  $\text{CH}_X=\text{CH}-\text{CH}=\text{O}$  (II, where X = Cl or Br), which are of great importance for further syntheses (Ref 3). It was found that in treating the compounds (I) with aqueous sodium carbonate solutions the corresponding acrolein (II) is formed.  $\beta$ -Halogen acroleins are colorless liquids, which assume a yellow color on standing and very strongly irritate the mucous membrane of the eyes and nose. When exposed to light they decompose already at room temperature under development of hydrogen halide. They are more stable at low temperatures in the presence of traces of hydroquinone or in the solution of an inert solvent. As was expected,  $\beta$ -halogen acroleins form pyrazole with semicarbazide or hydrazine, 1-arylpyrazoles with arylhydrazines, isoxazole with hydroxyl amine, 2-amino pyrimidine with guanidine, etc. The reactions take place over

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intermediate stages, in which the semicarbazones and aryl hydrazones of  $\beta$ -halogen acroleins could be isolated. On the basis of the synthesis of pyrazole the earlier suggested succession of reactions in the synthesis of heterocyclic compounds from the derivatives of  $\beta,\beta$ -dihalogen propionaldehydes may be considered to be an established fact, in so far as the authors succeeded in separating their semicarbazones and as far as the possibility of their easy transition into the semicarbazones of the  $\beta$ -halogen acroleins could be shown (Refs 1,8) (Scheme). Thus,  $\beta$ -chloro- and  $\beta$ -bromo acrolein in individual state were synthesized and characterized for the first time by the separation of hydrogen halide from the  $\beta,\beta$ -dihalogen propionaldehydes. The oxidation of  $\beta$ -halogen acroleins with silver oxide leads to the trans- $\beta$ -halogen acrylic acids. There are 13 references, 7 of which are Soviet.

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SUBMITTED: January 21, 1958

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AUTHORS:

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TITLE:

Investigation in the Series of Cyclobutanedicarboxylic Acids. I. The Study of the Structure of "Thesinic" Acid

PERIODICAL:

Zhurnal obshchey khimii, 1960, Vol 30, Nr 2, pp 484-488 (USSR)

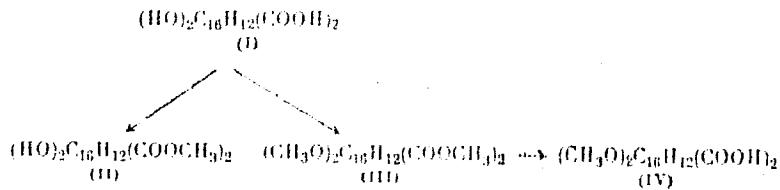
ABSTRACT:

In the previous paper (ZhOKh, 30, 670, abstract 77917) it was shown that the alkaloid thesin  $C_{34}H_{42}O_6N_2$  which was isolated from the plant Thesium minkwitzianum is an ester of d- "isoretronecanol" and a dibasic acid  $C_{18}H_{36}O_4$ . "Thesinic" acid has the following composition (I):

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butanedicarboxylic Acids. I

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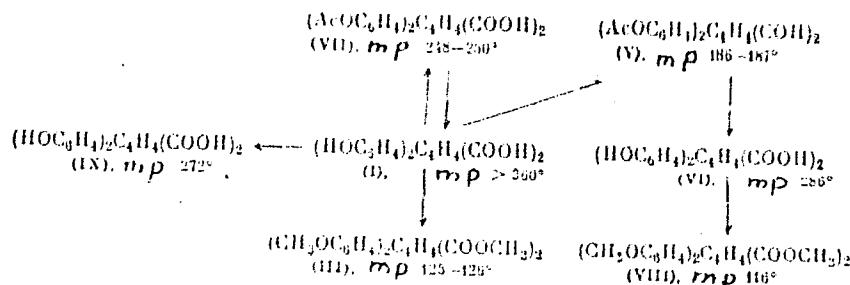


Methylation of I (1 g) with dimethyl sulfate (1.2 g) in the presence of 2 moles of  $\text{NaHCO}_3$  forms dimethyl thesinate (II) (0.5 g) (mp 183-184°). The above methylation (I, 3 g; dimethyl sulfate, 6.5 g), in the presence of 4 moles of NaOH, forms dimethyl dimethoxythesinate (III) (2.95 g) (mp 125-126°). Hydrolysis of III (3 g) with alcoholic alkali forms IV (1.8 g) (mp 250-251°). *p*-Hydroxycinnamic acid after a 20-hr exposure to direct sunlight forms "thesinic" acid, in 56% yield:

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V (0.8 g) was formed from I (1 g) on heating with acetic anhydride (5 ml) in the presence of sodium acetate (0.6 g) ( $mp$  186-187°). Hydrolysis of V with a  $\text{Na}_2\text{CO}_3$  solution forms VI ( $mp$  286). Acetylation of I (0.5 g) with glacial acetic acid (5 ml) and acetic anhydride (0.5 g) without sodium acetate forms VII (0.4 g) ( $mp$  248-250°). VI (0.3 g) with dimethyl sulfate (0.65 g), in the presence of  $\text{NaOH}$ , forms

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VIII (0.2 g) (mp 116°). IX(0.55 g) was obtained by fusion  
of I (1 g) with KOH (3 g) (mp 272°). There are  
5 references, 2 Soviet, 2 German, 1 U.S. The U.S.  
reference is: A. Mustafa, Chem. Revs., 51, 1 (1952).

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SUBMITTED: February 10, 1959

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